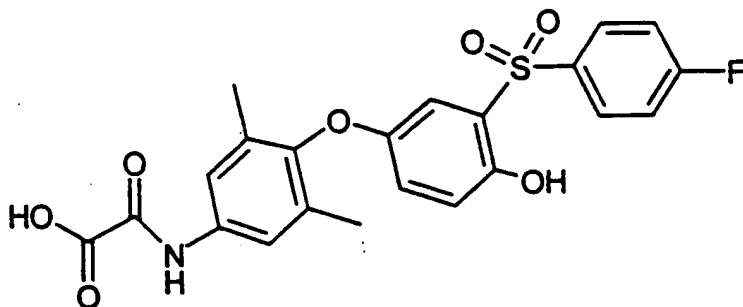


**EXHIBIT**

# Substance Sheet

Labjournal ID: U-0282-85-B  
Date:  
Department: RES/MCD/MCDCHEM  
Chem. Code:  
US Sample No:

Chemists: KUKKOLA, PAIVI  
WANG, HUA

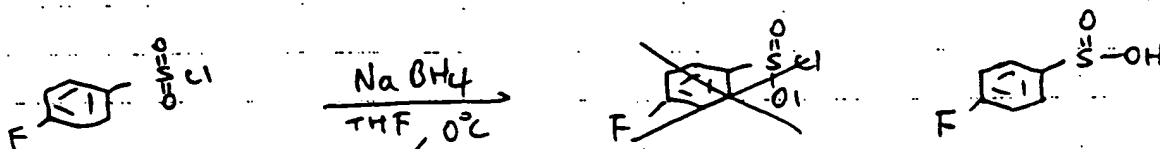


MW Eff.: 459.45  
Factor f: 1.000  
MF Subst.: C<sub>22</sub>H<sub>18</sub>FNO<sub>7</sub>S  
MW Subst.: 459.45  
Initial Amount: 50 mg

Subst./Stereo State: Single compd - achiral  
Struct. Assign.: Compatible with analytical data  
Approval Status: APPROVED  
Synthesis Sheet: Yes  
End Product: Yes  
Multi.Par / CChem.: No  
To be sent to LFU/NCA: No

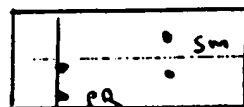
<sup>1</sup>H-NMR: Yes ; self service  
MS: 2919-  
IR: 2919-

Solvent System: DMSO  
Known in Lit.: No;  
To be tested on: LPM; 4 mg, R. Steel  
Clinical Codes:  
Comment Batch: CHN; 100% purity by HPLC



	Amount	mmol	eq
4-fluorobenzenesulfonyl chloride (194.61)	500 mg	2.57	1.0
Sodium borohydride (37.83)	0.49 g	12.85	5.0
THF	20 ml		

4-fluorobenzenesulfonyl chloride was dissolved in THF and NaBH<sub>4</sub> was added in portions with stirring at 0°C. The rxn was stirred at 0°C for 1 h, then RT for 5 h.

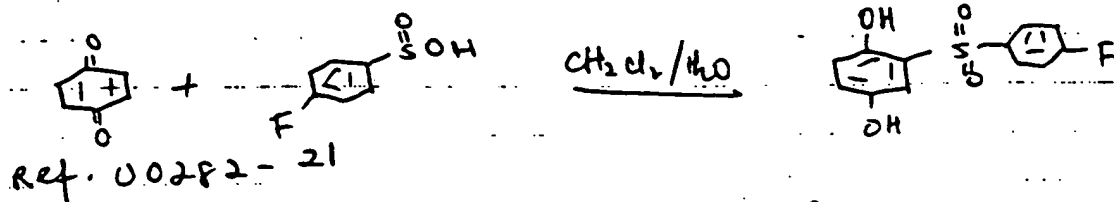


hex/EtOAc 3:2

After removal of THF, H<sub>2</sub>O was added and mixture was acidified by the addition of HCl (6N) dropwise at 0°C. Extracted w/ EtOAc. Dried over Na<sub>2</sub>SO<sub>4</sub> and concentrated to give 0.2-67 as a white solid. 330 mg

MS: (M+)<sup>+</sup> was detected

*Handwritten signature*



U0282-67 (160)

 $\text{CH}_2\text{Cl}_2$ 

1,4-benzoquinone (108.10)

 $\text{H}_2\text{O}$ 

Amount	mmol	g
320 mg	2.0	1.05
6 mL		
206 mg	1.9	1.0
4 mL		

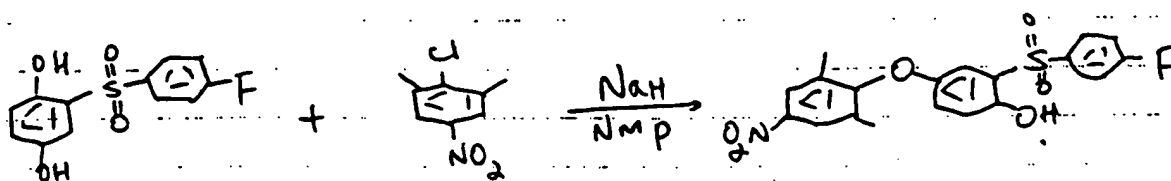
~~To a solution of U0282-67 in  $\text{CH}_2\text{Cl}_2$ , was added 1,4-benzoquinone~~

To a solution of 1,4-benzoquinone in  $\text{CH}_2\text{Cl}_2$ , was added a solution of U0282-67 in  $\text{H}_2\text{O}$ . The solution was stirred at RT. After 10 min, precipitation occurred. The suspension was stirred at RT for 3 h.

It was filtered. The solid was collected. NB U0282-70 200 mg as a off white solid

MS:  $\text{AcN}/\text{H}_2\text{O}/\text{NH}_4\text{OH}$   
 $(2M-1)^+$  was detected

Heun



Ref 00282-75

00282-70 (268)

NaH (24, 60%)

NB 2694-12 (185.6)

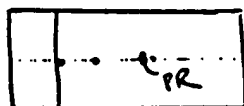
NMP

Amount	mmol	g
200mg	0.75	1.0
99mg	2.46	3.3
167mg	0.90	1.2
5ml		

To a suspension of NaH in NMP at 0°C, was added 00282-70. The suspension was stirred at RT for 30 min. NB 2694-12 was added. The black suspension was stirred at 100°C for 3h.

It was quenched w/ H<sub>2</sub>O and extracted w/ Et<sub>2</sub>O (x3). The organic layer was washed w/ brine, dried over Na<sub>2</sub>SO<sub>4</sub> and concentrated to give 00282-74 as a yellow solid.

MS: (M-1)<sup>+</sup> was detected (416)

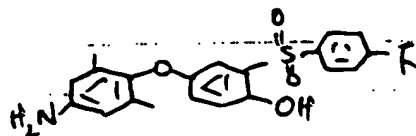
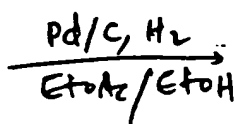
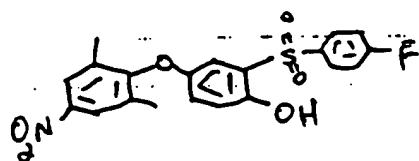


hex/EtOAc 3:2

The crude was purified w/ flash chromatography (hex/EtOAc 3:2) to give 00282-74A. 246mg as a white solid.

<sup>1</sup>H NMR (CDCl<sub>3</sub>) of 00282-74A: reasonably clean

Huan



Ref. U0282-38

U0282-74A

Pd/C

EtOH/EtOAc

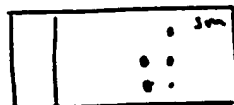
Amount mmol of

240 mg

24 mg

10 ml / 10 ml

The rxn was stirred at RT & under H<sub>2</sub> (1 atm) for 17 h.

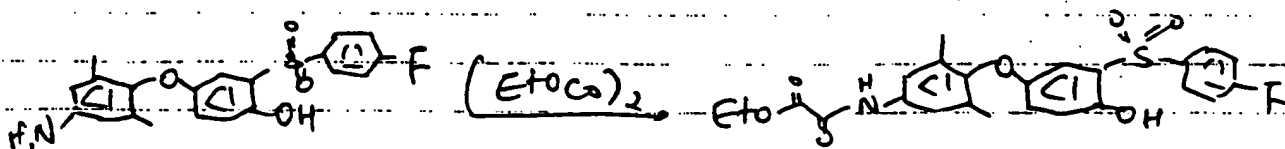
H<sub>2</sub>/EtOAc 2:3

Small amount of starting material still present. It was stirred under H<sub>2</sub> balloon at RT for 4 h. It was completed by TLC. It was filtered through celite. The filtrate was concentrated to give U0282-80 190 mg as a solid.

<sup>1</sup>H NMR (CDCl<sub>3</sub>)

OK

H<sub>2</sub>O



Ref. U0282-38

U0282-80

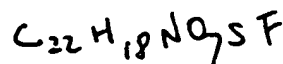
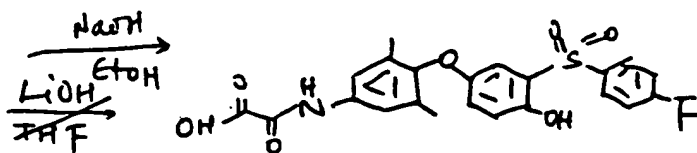
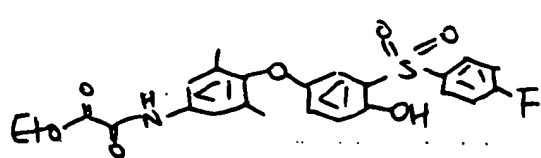
(EtOCO)<sub>2</sub>

Amount	mmol	g
1.90mg		
2ml		

The rxn was stirred at 180°C for 3h. The solvents were removed by N<sub>2</sub> stream. Chromatographed on silica (hex/EtOAc 2:1) to give U0282-83A as a yellowish foam 195mg.

<sup>1</sup>H NMR (CDCl<sub>3</sub>): OK. trace amount of impurities.

Hua ~~~~~



Ref 00212-45

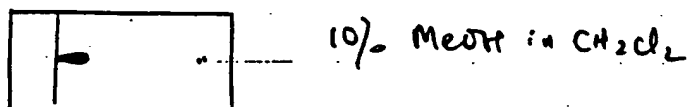
00212-83 (487.5)

NaOH (1M)

EtOH

Amount	mmol	eq
170 mg	0.35	1.0
1.05 ml	1.05	3.0
3 ml		

The rxn was stirred at RT for 2 h.

10% MeOH in  $CH_2Cl_2$ 

It was quenched w/ HCl (1N, 1.5 ml). Extracted w/ EtOAc. The organ was washed w/ brine. Dried over  $Na_2SO_4$  and concentrated to give 00212-83A as a foam.

00212-83A was triturated w/ Et<sub>2</sub>O/hexane. Dried to give 00212-85A 75 mg as a white solid.

<sup>1</sup>H NMR of 00212-85A: same ether.

00212-85A was dried in vacuum over at 50°C for h. to give 00212-85 57 mg as a white solid.

<sup>1</sup>H NMR ( $CDCl_3$ ): OK.

MS: (m-1)<sup>-</sup> detected.

HN: OK.

50 mg sample sent out.

Heuer



Sheet1

	nM	Reading1	Reading2	Mean	Activity(%)	Estimate	Hill Coefficient	IC-50
NO ENZ		494.00	494.00	494.00	0.00		0.84427425	0.167190443
NO INH		17268.00	17268.00	17268.00	100.00			
	0.010	15583.00	15583.00	15583.00	89.95	91.51		
	0.030	14660.00	14660.00	14660.00	84.45	81.01		
	0.060	11483.00	11483.00	11483.00	65.51	70.37		
	0.100	11450.00	11450.00	11450.00	65.32	60.68		
	0.150	9689.00	9689.00	9689.00	54.82	52.29		
	0.200	9572.00	9572.00	9572.00	54.12	46.23		
	0.250	7859.00	7859.00	7859.00	43.91	41.59		
	0.300	4881.00	4881.00	4881.00	26.15	37.90		

Concentrations	% Inhibition	Calculated
0.010	10.045	8.49
0.030	15.548	18.99
0.060	34.488	29.63
0.100	34.685	39.32
0.150	45.183	47.71
0.200	45.881	53.77
0.250	56.093	58.41
0.300	73.846	62.10

IC50

*Jameson Whelan*

*[Large handwritten X mark]*

Read and understood by me

*Chibola*

Date